

EXHIBIT 26

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Sent: 12/20/2017 8:06:58 PM
Subject: NOTIFICATION: UNANNOUNCED DEA Inspection at the Kentucky Distribution Center (KDC) 20 Dec 2017
Attachments: Regulatory Inspection – Importer Exporter Check List.pdf

Hi, All,

Today, KDC had an unannounced DEA inspection. The inspection was a routine inspection on KDC's importer and exporter DEA Registrations.

Overall re-cap – **zero (0) observations/opportunities**. The Diversion Investigators, Carl and Morgan, had nothing but positive feedback.

I have attached the 'Regulatory Inspection – Importer Exporter Check List' that the DEA is using for Importer and Exporter inspections. This is a handy check list that I'm asking you all to share with your contacts. We went line by line with our investigators at KDC today, so if you have all of this documentation handy and updated, your future inspections should go smoothly.

As an FYI, there was a discussion on the Suspicious Order Monitoring process. The DEA confirmed that although the Customer Service group has responsibilities over this process, any deficiencies against this process would be an observation for KDC.

Thanks to everyone that supported...Christian, Mark, Kevin, Christina, Josh, Morgan, Jon, Belinda and Brad! Curve ID 1001617 was initiated for this inspection.

Thanks,

KDC Team

INSPECTION COMMUNICATION:

INSPECTING REGULATORY BODY/BODIES	DEA Attendee (s): B. Morgan Freeman, Diversion Investigator Carl Maskew, Diversion Investigator
SCOPE OF INSPECTION/PRODUCTS INVOLVED & STAGE OF MANUFACTURE (e.g. API, fill/finish, packaging, etc.)	3 Year Inspection on Importer/Exporter
TYPE OF INSPECTION (e.g. GMP, Pre-Approval, Directed)	3 Year Inspection on Importer/Exporter
DATE (anticipated or actual start date)	20 Dec 2017 Start Time – 9:30 a.m. End Time – 2:15 p.m.
LOCATION OF INSPECTION	Kentucky Distribution Center (KDC)
ISSUES IDENTIFIED, if available	No issues; zero (0) observations



KEY CONTACTS (e.g. Quality Head, Inspection host/lead)

Brad Hummel, Director Distribution Operations, KDC

Rob Helfrick, Sr. Manager Quality and Compliance, KDC

Mark Hood, Operations Supervisor, KDC

Christian Williamson, Inventory Control and Analytics Manager, KDC

Kevin Pendigraph, Allied Universal Security Manager, KDC

Jon Ostermann, Site Facilities Manager, KDC

Belinda Corum, Controlled Substance Specialist, KDC

3 yrs - oct - sept - fiscal yrs

REGULATORY INSPECTION - IMPORTER/EXPORTER

The following information is required during the Regulatory Inspection of your firm. Please provide this information/documentation to investigators during their on-site inspection.

✓ 1.	List of the firm's Executive Officers (Full Name, Date of Birth, Social Security Number, Position/Title)	
✓ 2.	List of the firm's Corporate Officers (Full Name, Date of Birth, Social Security Number, Position/Title)	
✓ 3.	List of all Employees (Full Name & Position/Title)	MH
✓ 4.	List of all Employee who have access to Controlled Substances (Breakdown by: Key Access, Alarm Code Access, or Both) (Full Name, Date of Birth, Social Security Number, Position/Title)	MH
✓ 5.	List of all Controlled Substances handled by the firm (Drug Name, Product Size, Product Strength, and Product Form)	CS import CS export CW
6.	List all Listed Chemicals handled by the firm (Name, Product Size, Product Strength, and Product Form) - Note: Pseudoephedrine and Ephedrine are Listed Chemicals.	n/a
7.	List of all Licenses, Permits, & Registrations issued to firm (Issuing Agency, Type of Registration, Issue Date, Expiration Date)	MH
8.	Initial Inventories	n/a
9.	Biennial Inventory	Done
10.	Power of Attorney Forms pertaining to Schedule II Controlled Substances	
11.	Supplier List (Name, Address, Phone, & DEA #)	Export Supplier
12.	Customer List (Name, Address, Phone, & DEA #)	n/a
13.	Map/Floorplan of facility	Done - SA
14.	Alarm Information (Description, Security Contract, Schematic, Maintenance/Test Records)	Done - SA
✓ 15.	All written policies or procedures pertaining to Controlled Substances (i.e. Standard Operating Procedures) - Please include Due Diligence for New Customers and New Patients	
16.	Records required for designated Audit Period (PLEASE PROVIDE A COMPUTER PRINTOUT OF RECEIPTS AND DISTRIBUTIONS & DIS WILL RANDOMLY	n/a

Only importer

	<p>SELECT RECORDS FROM BELOW TO RECONCILE WITH PRINTOUT)</p> <ul style="list-style-type: none"> • DEA-222 Forms (if handle CII) • Records of Receipt of Controlled Substance • Records of Sales/Distribution of Controlled Substances • Records of Returned Controlled Substances from Customers • Records of Returned Controlled Substances to Distributors • Records of Damaged Controlled Substances (DEA-42 if applicable) • Records of Theft or Loss of Controlled Substances (DEA-106) • Controlled Substance Import/Export Declarations (DEA-236) • Application for Permit to Import Controlled Substances for Domestic &/or Scientific Purposes (DEA-357) 	n/s
17.	Subsidiaries (Name, Address, Phone, DEA #) - If applicable	Pending
18.	Parent Companies (Name, Address, Phone, DEA #) - If applicable	
19.	Drug Test Company (Name, Address, Phone)	
20.	Background Check Company (Name, Address, Phone)	
21.	Export authorizations by country of destination	n/s
22.	Articles of Incorporation	

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